## Listing of claims

Please amend claims 8, 47, 67-70, 72, and 79 as shown below. Please cancel claims 10, 46, 48, 60, 66, 71, 76-78 and withdrawn claims 54-58, 61-65 and 73-75.

- 1-7. (canceled)
- 8. (currently amended) The stent of claim 4647 comprising:
- a radially self-expanding tubular shaped member having first and second ends; a walled surface disposed between said first and second ends;

said walled surface comprising a plurality of substantially parallel pairs of monofilaments; said substantially parallel pairs of monofilaments woven in a helical shape such that substantially one-half of said substantially parallel pairs of monofilaments are wound clockwise in the longitudinal direction and one-half of said substantially parallel pairs of monofilaments are wound counterclockwise in the longitudinal direction such that an alternating, over-under plait of said substantially parallel pairs of monofilaments results; said monofilaments consisting essentially of the blend of two bioresorbable, bio-compatible homopolymers.

- 9. (previously presented) The stent of claim 8, comprising approximately twenty-four substantially parallel pairs of monofilaments.
  - 10. (canceled)
- 11. (previously presented) The stent of claim 8, wherein said polymer blend possesses a tensile strength in the range of approximately 40,000 psi to 120,000 psi.
- 12. (previously presented) The stent of claim 8, wherein said polymer blend possesses a tensile modulus in the range of approximately 400,000 psi to 2,000,000 psi.

- 13. (previously presented) The bioresorbable stent of claim 8, wherein said stent has a compressed first diameter of between approximately 6 mm and 10 mm and a second non-compressed diameter of between approximately 12 mm and 18 mm.
- 14. (previously presented) The bioresorbable stent of claim 8 wherein said woven monofilaments have a crossing angle of between approximately 100 degrees and 150 degrees in the non-compressed resting state.
  - 15-45. (canceled)
  - 46. (canceled)
- 47. (currently amended) The stent of claim 4681 comprising:
  a walled surface disposed between said first and second ends;
  said walled surface comprising a helical shape of woven monofilaments
  consisting essentially of comprising the blend of two bioresorbable, bio-compatible homopolymers.
  - 48. (canceled)
- 49. (previously presented) The stent of claim 47, wherein said walled surface has approximately 30 monofilaments.
- 50. (previously presented) The stent of claim 47, wherein said polymer blend possesses a tensile strength in the range of approximately 40,000 psi to 120,000 psi.
- 51. (previously presented) The stent of claim 47, wherein said polymer blend possesses a tensile modulus in the range of approximately 400,000 psi to 2,000,000 psi.
- 52. (previously presented) The stent of claim 47, wherein said stent has a compressed first diameter of between approximately 6 mm and 10 mm and a second non-compressed diameter of between approximately 12 mm and 18 mm.

53. (previously presented) The stent of claim 47, wherein said woven monofilaments have a crossing angle of between approximately 100 degrees and 150 degrees in the non-compressed resting state.

54-58. (canceled)

- 59. (previously presented) The stent of claim 46, wherein the stent is a urethral stent.
  - 60. (canceled)
  - 61-66. (canceled)
- 67. (currently amended) The stent of claim 6672, wherein poly-L-lactide and poly-ε-caprolactone are present at a ratio between approximately 80:20 and 99:1.
- 68. (currently amended) The stent of claim 6780, wherein the ratio is approximately 90:10.
- 69. (currently amended) The stent of claim 6679 wherein the poly-L-lactide has a molecular weight of approximately 450,000 daltons or greater.
- 70. (currently amended) The stent of claim 6979 wherein the poly-L-lactide has a molecular weight of approximately 750,000 daltons or greater.
  - 71. (canceled)
- 72. (currently amended) A bioresorbable, self-expanding stent having a non-compressed diameter of between approximately 12 millimeters and 18 millimeters comprising

a radially self-expanding tubular-shaped bioresorbable member having first and second ends, a walled surface disposed between said first and second ends;

said walled surface comprising a plurality of substantially parallel pairs of monofilaments; said substantially parallel pairs of monofilaments woven in a helical shape such that substantially one-half of said substantially parallel pairs of monofilaments are wound clockwise in the longitudinal direction and one-half of said substantially parallel pairs of monofilaments are wound counterclockwise in the longitudinal direction such that an alternating, over-under plait of said substantially parallel pairs of monofilaments results; said monofilaments consisting essentially of poly-L-lactide and poly-ε-caprolactone, The stent of claim 71 wherein the poly-ε-caprolactone has a molecular weight of approximately 200,000 daltons or greater.

73-75. (canceled)

76-78. (canceled)

- 79. (currently amended) A<u>The</u> bioresorbable, self-expanding stent of claim 81 comprising a tubular shaped bioresorbable member having first and second ends, said bioresorbable member comprising poly-L-lactide and poly-ε-caprolactone homopolymers, the stent having a non-compressed diameter of between approximately 12 millimeters and 18 millimeters.
- 80. (previously presented) The stent of claim 79, wherein poly-L-lactide and poly-ε-caprolactone are present at a ratio between approximately 80:20 and 99:1.
- 81. (previously presented) A bioresorbable, self-expanding stent comprising a tubular-shaped bioresorbable member having first and second ends, said bioresorbable member comprising a blend of at least two bioresorbable, bio-compatible homopolymers, wherein one of the two homopolymers is poly-ε-caprolactone having a molecular weight of approximately 200,000 daltons or greater, the stent having a non-compressed diameter of between approximately 12 millimeters and 18 millimeters.